What is claimed is:

- 1 1. A composite electrolyte for use in electrochemical fuel cells, comprising:
- 2 (i) an inorganic cation exchange material;
- 3 (ii) a silica-based material; and
- 4 (iii) a proton conducting polymer-based material, wherein the inorganic cation
- 5 exchange material comprises about 0.1 wt% to about 99 wt% of the composite
- 6 electrolyte.
- 1 2. The composite electrolyte of claim 1, wherein the silica-based material comprises
- 2 about 0.1 wt% to about 70 wt%, and the proton conducting polymer-based material
- 3 comprises about 0.1 wt% to 99.9 wt% of the composite electrolyte.
- 1 3 The composite electrolyte of claim 1 wherein the inorganic cation exchange material
- 2 is selected from the group consisting of clay, zeolite, hydrous oxide, and inorganic salt.
- 1 4. The composite electrolyte of claim 3, wherein the clay includes an aluminosilicate-
- 2 based exchange material selected from the group consisting of montmorillonite, kaolinite,
- 3 vermiculite, smectite, hectorite, mica, bentonite, nontronite, beidellite, volkonskoite,
- 4 saponite, magadite, kenyaite, zeolite, alumina, rutile.
- 1 5. The composite material of claim 3, wherein the clay is modified to make it more
- 2 compatible with organic matrices, a clay modification including exfoliation which helps to
- 3 separate platelets of inorganic substance.
- 1 6. The composite electrolyte of claim 3, wherein the clay includes a modified
- 2 montmorillonite consisting of montmorillonite treated with a modifier material selected from
- a group consisting of aminododecanoic acid, trimethyl stearate ammonium, octadecylamine,
- 4 and methyl dihydroxy hydrogenated tallow ammonium.
- 1 7. The composite electrolyte of claim 1 wherein the inorganic cation exchange material
- 2 comprises about 0.1 wt% to about 30 wt%, the silica-based material comprises about 0.1 wt%
- 3 to about 30 wt%, and the proton conducting polymer-based material comprises about 40 wt%
- 4 to 99.9 wt% of the composite electrolyte.

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- 1 8. The composite electrolyte of claim 1 wherein the proton conducting polymer-based
- 2 material has a linear, branched, or network morphology.
- 1 9. The composite electrolyte of claim 1 wherein the proton conducting polymer-based
- 2 material includes material selected from the group consisting of
- 3 acrylonitrile/butadiene/stryene rubber (ABS), styrene butadiene/acrylate/acetate polymer
- 4 blends, epoxides, and a thermoplastic material.
- 1 10. The composite electrolyte of claim 9 wherein the thermoplatic material is selected
- 2 from a group consisting polypropylene, polycarbonate, polystyrene, polyethylene, polyaryl
- 3 ethers sulfones, poly aryl ether ketone, and polysulfones.
- 1 11. The composite electrolyte of claim 1 wherein the proton conducting polymer-based
- 2 material has a functional group element selected from a group consisting of sulfonate,
- 3 phosphate, carbonate, amide, and imide, wherein each such functional group element has
- 4 proton conducting capabilities.
- 1 12. The composite electrolyte of claim 1, further comprising an additive selected from a
- 2 group consisting of preservative, thixotropy and viscosity control agent, crosslinking agent,
- 3 conditioner, plasticizer, water control agent, and proton conducting material.
- 1 13. The composite electrolyte of claim 1 wherein the inorganic cation exchange material,
- 2 the silica-based material and the proton conducting polymer-based material comprise 90 wt
- 3 % or more of the solids content of the composite electrolyte.
- 1 14. The composite electrolyte of claim 1 wherein the composite electrolyte when
- 2 measured in the substantially dried state consists essentially of the inorganic cation exchange
- 3 material, the silica-based material and the proton conducting polymer-based material.
- 1 15. The composite electrolyte of claim 1 wherein the composite electrolyte has a proton
- 2 conductivity of about 0.05 S/cm or higher

16. The composite electrolyte of claim 1 wherein the silica-based material includes materials containing one or more of silica, silicate, and silicate having an organic element. 2 The composite electrolyte of claim 1 wherein the silica-based material is either 17. 1 colloidal silica containing discrete spheres of silica or tetraethylorthosilicate. 2 An electrochemical fuel cell, comprising: 18. an anode; (i) 2 a cathode; (ii) 3 a fuel supply to the anode; (iii) 4 an oxidant supply to the cathode; and (iv) 5 (v) a composite electrolyte positioned between the anode and cathode and 6 including 7 (a) an inorganic cation material, 8 (b) a silica-based binder, and 9 (c) a polymer-based binder, 10 wherein the inorganic cation exchange material comprises about 0.1 wt% to about 99 11 wt%, of the composite electrolyte. 12 The electrochemical fuel cell of claim 18 wherein the silica-based material comprises 19. about 0.1 wt% to about 70 wt%, and the proton conducting polymer-based material 2 comprises about 0.1 wt% to 99.9 wt% of the composite electrolyte. 3 The electrochemical fuel cell of claim 18 wherein the inorganic cation exchange material comprises about 0.1 wt% to about 30 wt%, the silica-based material comprises about 2 0.1 wt% to about 30 wt%, and the proton conducting polymer-based material comprises 3 about 40 wt% to 99.9 wt% of the composite electrolyte. 4 The electrochemical fuel cell of claim 18 wherein the inorganic cation exchange 21. material is selected from the group consisting of clay, zeolite, hydrous oxide, and inorganic 2 salt. 3 The electrochemical fuel cell of claim 21 wherein the clay includes an 22. aluminosilicate-based exchange material selected from the group consisting of 2 73276.4.17 8/19/2003 Page 18 of 23 372584-337162 Taft et al.

- 3 montmorillonite, kaolinite, vermiculite, smectite, hectorite, mica, bentonite, nontronite,
- 4 beidellite, volkonskoite, saponite, magadite, kenyaite, zeolite, alumina, and rutile.
- 1 23. The electrochemical fuel cell of claim 21, wherein the clay is modified to make it
- 2 more compatible with organic matrices, a clay modification including exfoliation which helps
- 3 to separate platelets of inorganic substance.
- 1 24. The electrochemical fuel cell of claim 21, wherein the clay includes a modified
- 2 montmorillonite consisting of montmorillonite treated with a modifier material selected from
- 3 a group consisting of aminododecanoic acid, trimethyl stearate ammonium, octadecylamine,
- 4 and methyl dihydroxy hydrogenated tallow ammonium.
- 1 25. The electrochemical fuel cell of claim 18 wherein the polymer-based material has
- 2 linear, branched, or network morphology.
- 1 26. The electrochemical fuel cell of claim 18 wherein the polymer-based material
- 2 includes material selected from the group consisting of acrylonitrile/butadiene/stryene rubber
- 3 (ABS), styrene butadiene/acrylate/acetate polymer blends, epoxides, polypropylene,
- 4 polycarbonate, polystyrene, polyethylene, polyaryl ethers, and polysulfones.
- 1 27. The electrochemical fuel cell of claim 18 wherein the inorganic cation exchange
- 2 material, the silica-based material and the polymer-based material comprise 90 wt % or more
- of the solids content of the composite electrolyte.
- 1 28. The electrochemical fuel cell of claim 18 wherein the composite electrolyte when
- 2 measured in the substantially dried state consists essentially of the inorganic cation exchange
- 3 material, the silica-based material and the polymer-based material.
- 1 29. The electrochemical fuel cell of claim 18 wherein the composite electrolyte has a
- 2 proton conductivity of about 0.05 S/cm or higher.
- 1 30. A method of fabricating a composite electrolyte for use in an electrochemical fuel
- 2 cell, comprising:

- applying onto a surface of a substrate a viscous liquid composition of (a) an (i) 3 inorganic cation exchange material, (b) silica-based material, (c) a polymer-4 based material, and (d) a solvent-dispersant; 5 spreading the viscous liquid composition to form a uniform thickness layer on (ii) 6 the substrate; and 7 allowing the solvent to evaporate from the viscous liquid composition to yield (iii) 8 the composite electrolyte, 9 wherein the inorganic cation exchange material comprises about 0.1 wt% to 10
- 1 31. The method of claim 30, wherein the silica-based material comprises about 0.1 wt%
- 2 to about 70 wt%, and the polymer-based material comprises about 0.1 wt% to 99.9 wt% of
- 3 the composite electrolyte.
- 1 32. The method of claim 30 wherein step (ii) includes drawing the viscous liquid

about 99 wt% of the composite electrolyte.

- 2 composition through a doctor blade assembly.
- 1 33. The method of claim 30 wherein step (iii) includes heating the viscous liquid
- 2 composition.

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- 1 34. The method of claim 30 wherein the inorganic cation exchange material comprises
- 2 about 0.1 wt% to about 30 wt%, the silica-based material comprises about 0.1 wt% to about
- 3 15 wt%, and the polymer-based material comprises about 40 wt% to 99 wt% of the composite
- 4 electrolyte.
- 1 35. The method of claim 19 wherein the inorganic cation exchange material is selected
- 2 from the group consisting of clay, zeolite, hydrous oxide, inorganic and salt.
- The method of claim 35 wherein the clay includes an aluminosilicate-based exchange
- 2 material selected from the group consisting of montmorillonite, kaolinite, vermiculite,
- 3 smectite, hectorite, mica, bentonite, nontronite, beidellite, volkonskoite, saponite, magadite,
- 4 kenyaite, zeolite, alumina, and rutile.

- 1 37. The method of claim 35, wherein the clay is modified to make it more compatible
- 2 with organic matrices, a clay modification including exfoliation which helps to separate
- 3 platelets of inorganic substance.
- 1 38. The method of claim 35, wherein the clay includes a modified montmorillonite
- 2 consisting of montmorillonite treated with a modifier material selected from a group
- 3 consisting of aminododecanoic acid, trimethyl stearate ammonium, octadecylamine, and
- 4 methyl dihydroxy hydrogenated tallow ammonium.
- 1 39. The method of claim 30 wherein the polymer-based material has a linear, branched, or
- 2 netted morphology.
- 1 40. The method of claim 30 wherein the polymer-based material includes one of
- 2 acrylonitrile/butadiene/stryene rubber (ABS), styrene butadiene/acrylate/acetate polymer
- 3 blends, epoxides, polypropylene, polycarbonate, polystyrene, polyethylene, polyaryl ethers,
- 4 and polysulfones.
- 1 41. The method of claim 30 wherein the solvent-dispersant comprises water, N-methyl
- 2 pyrrolidinone, dimethyl sulfoxide, dimethyl acidimide, and dimethylformamide.
- 1 42. The method of claim 30 wherein the inorganic cation exchange material, the silica-
- 2 based material and the polymer-based material comprise 90 wt % or more of the solids
- 3 content of the composite electrolyte.
- 1 43. The method of claim 30 wherein the composite electrolyte when measured in the
- 2 substantially dried state consists essentially of the inorganic cation exchange material, the
- 3 silica-based material and the polymer-based material.
- 1 44. The method of claim 19 wherein the composite electrolyte has a proton conductivity
- of about 0.05 S/cm or higher.
- 1 45. A method for producing a composite membrane, comprising:

- 2 (i) grinding a sulfonated polyether ether ketone (SPEEK) and clay mixture and
- dissolving the mixture in a distilled dimethylformamide (DMF) to form a solution;
- 4 (ii) heating the solution until it thickens and attains a casting consistency;
- 5 (iii) degassing the solution in a vacuum oven;
- 6 (iv) casting the solution into a film on a glass surface using a doctor blade;
- 7 (v) curing the film; and
- 8 (vi) peeling the film from the glass.
- 1 46. The method of claim 45, wherein the dissolving in step (i) is performed by stirring
- 2 for about 2 hours using a magnetic stir bar.
- 1 47. The method of claim 45, wherein the solution is stirred while heated, and wherein the
- 2 DMF evaporates.
- 1 48. The method of claim 45, wherein the film is about 60 μm thick.
- 1 49. The method of claim 45, wherein the curing includes,
- 2 (a) annealing the film in a convection oven, and
- 3 (b) maintaining the film in a vacuum for a predetermined time period at a
- 4 predetermined temperature.
- 1 50. The method of claim 45, further comprising:
- 2 storing the film in ultra-pure water until it is ready for use.